

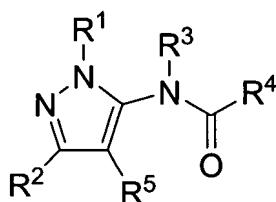
**AMENDMENTS TO THE CLAIMS**

Please cancel Claims 1-20 and insert therefor Claims 21-35 as follow. This listing of claims will replace all prior versions, and listings, of claims in the application.

**Listing of Claims:**

1-20. (Canceled)

21. (New) A compound of the formula I:



I

wherein:

R<sup>1</sup> is selected from the group consisting of:

- (1) hydrogen,
- (2) C<sub>1</sub>-6alkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl,
- (3) C<sub>3</sub>-7cycloalkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl, and
- (4) phenyl, which is unsubstituted or substituted with one or more substituents independently selected from:
  - (a) -C<sub>1</sub>-6alkyl,
  - (b) -O-C<sub>1</sub>-6alkyl,
  - (c) halo,
  - (d) hydroxy,
  - (e) trifluoromethyl,
  - (f) -OCF<sub>3</sub>,
  - (g) -CO<sub>2</sub>R<sup>9</sup>,

wherein R<sup>9</sup> is independently selected from:

- (i) hydrogen,
- (ii) -C<sub>1</sub>-6alkyl, which is unsubstituted or substituted with 1-6 fluoro,
- (iii) benzyl, and

- (iv) phenyl,
  - (h) -NR<sup>10</sup>R<sup>11</sup>,  
wherein R<sup>10</sup> and R<sup>11</sup> are independently selected from:
    - (i) hydrogen,
    - (ii) -C<sub>1-6</sub>alkyl, which is unsubstituted or substituted with 1-6 fluoro,
    - (iii) -C<sub>5-6</sub>cycloalkyl,
    - (iv) benzyl,
    - (v) phenyl,
    - (vi) -S(O)<sub>2</sub>-C<sub>1-6</sub>alkyl,
    - (vii) -S(O)<sub>2</sub>-benzyl, and
    - (viii) -S(O)<sub>2</sub>-phenyl,
  - (i) -CONR<sup>10</sup>R<sup>11</sup>, and
  - (j) -NO<sub>2</sub>;
- (5) heterocycle, wherein heterocycle is selected from:  
benzoimidazolyl, benzimidazolonyl, benzofuranyl, benzofurazanyl,  
benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl,  
carbolinyl, cinnolinyl, furanyl, imidazolyl, indolinyl, indolyl, indolazinyll,  
indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl,  
naphthpyridinyl, oxadiazolyl, oxazolyl, oxazoline, isoxazoline, oxetanyl,  
pyranyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridopyridinyl, pyridazinyl, pyridyl,  
pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, tetrahydropyranlyl,  
tetrazolyl, tetrazolopyridyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidinyll,  
1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl, pyridin-2-onyl,  
pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzoimidazolyl,  
dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl,  
dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl,  
dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl,  
dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl,  
dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl,  
dihydrothienyl, dihydrotriazolyl, dihydroazetidinyll, methylenedioxybenzoyl,  
tetrahydrofuranyl, and tetrahydrothienyl, and N-oxides thereof, which is  
unsubstituted or substituted with one or more substituents independently  
selected from:
- (a) -C<sub>1-6</sub>alkyl,
  - (b) -O-C<sub>1-6</sub>alkyl,
  - (c) halo,

- (d) hydroxy,
- (e) phenyl,
- (f) trifluoromethyl,
- (g) -OCF<sub>3</sub>,
- (h) -CO<sub>2</sub>R<sup>9</sup>,
- (i) -NR<sup>10</sup>R<sup>11</sup>, and
- (j) -CONR<sup>10</sup>R<sup>11</sup>;

R<sup>2</sup> is phenyl;

R<sup>3</sup> is independently selected from the group consisting of:

- (1) hydrogen, and
- (2) C<sub>1</sub>-6alkyl;

R<sup>4</sup> is selected from the group consisting of:

- (1) C<sub>1</sub>-6alkyl, which is unsubstituted or substituted with halogen, hydroxyl, phenyl or heterocycle,
- (2) C<sub>3</sub>-7cycloalkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl, and
- (3) phenyl, which is unsubstituted or substituted with one or more substituents independently selected from:
  - (a) -C<sub>1</sub>-6alkyl,
  - (b) -O-C<sub>1</sub>-6alkyl,
  - (c) halo,
  - (d) hydroxy,
  - (e) trifluoromethyl,
  - (f) -OCF<sub>3</sub>,
  - (g) -CO<sub>2</sub>R<sup>9</sup>,
  - (h) -CN,
  - (i) -NR<sup>10</sup>R<sup>11</sup>,
  - (j) -CONR<sup>10</sup>R<sup>11</sup>, and
  - (k) -NO<sub>2</sub>;
- (4) heterocycle, wherein heterocycle is selected from:  
benzoimidazolyl, benzimidazolonyl, benzofuranyl, benzofurazanyl,  
benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl,  
carbolinyl, cinnolinyl, furanyl, imidazolyl, indolinyl, indolyl, indolazinyl,  
indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl,  
naphthpyridinyl, oxadiazolyl, oxazolyl, oxazoline, isoxazoline, oxetanyl,

pyranyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridopyridinyl, pyridazinyl, pyridyl, pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxaliny, tetrahydropyranyl, tetrazolyl, tetrazolopyridyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidyl, 1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl, pyridin-2-onyl, pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzoimidazolyl, dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl, dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl, dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl, dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl, dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl, dihydrothienyl, dihydrotriazolyl, dihydroazetidyl, methylenedioxybenzoyl, tetrahydrofuranyl, and tetrahydrothienyl, and N-oxides thereof, which is unsubstituted or substituted with one or more substituents independently selected from:

- (a) -C<sub>1-6</sub>alkyl,
- (b) -O-C<sub>1-6</sub>alkyl,
- (c) halo,
- (d) hydroxy,
- (e) phenyl,
- (f) trifluoromethyl,
- (g) -OCF<sub>3</sub>,
- (h) -CO<sub>2</sub>R<sup>9</sup>,
- (i) -NR<sup>10</sup>R<sup>11</sup>, and
- (j) -CONR<sup>10</sup>R<sup>11</sup>;

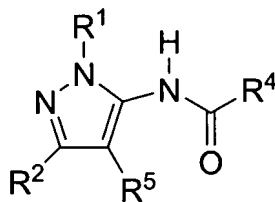
R<sup>5</sup> is independently selected from the group consisting of:

- (1) hydrogen,
- (2) C<sub>1-6</sub>alkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl,
- (3) C<sub>3-7</sub>cycloalkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl, and
- (4) phenyl, which is unsubstituted or substituted with one or more substituents independently selected from:
  - (a) -C<sub>1-6</sub>alkyl, which is unsubstituted or substituted with -NR<sup>10</sup>R<sup>11</sup>,
  - (b) -O-C<sub>1-6</sub>alkyl,
  - (c) halo,

- (d) hydroxy,
  - (e) trifluoromethyl,
  - (f) -OCF<sub>3</sub>;
  - (g) -CO<sub>2</sub>R<sup>9</sup>,
  - (h) -NR<sup>10</sup>R<sup>11</sup>,
  - (i) -C(O)NR<sup>10</sup>R<sup>11</sup>, and
  - (j) -NO<sub>2</sub>,
- (5) heterocycle, wherein heterocycle is selected from:  
benzoimidazolyl, benzimidazolonyl, benzofuranyl, benzofurazanyl,  
benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl,  
carbolinyl, cinnolinyl, furanyl, imidazolyl, indolinyl, indolyl, indolazinyll,  
indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl,  
naphthpyridinyl, oxadiazolyl, oxazolyl, oxazoline, isoxazoline, oxetanyl,  
pyranyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridopyridinyl, pyridazinyl, pyridyl,  
pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, tetrahydropyranyl,  
tetrazolyl, tetrazolopyridyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidinyll,  
1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl, pyridin-2-onyl,  
pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzoimidazolyl,  
dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl,  
dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl,  
dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl,  
dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl,  
dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl,  
dihydrothienyl, dihydrotriazolyl, dihydroazetidinyll, methylenedioxybenzoyl,  
tetrahydrofuranyl, and tetrahydrothienyl, and N-oxides thereof, which is  
unsubstituted or substituted with one or more substituents independently  
selected from:
- (a) -C<sub>1-6</sub>alkyl,
  - (b) -O-C<sub>1-6</sub>alkyl,
  - (c) halo,
  - (d) hydroxy,
  - (e) phenyl,
  - (f) trifluoromethyl,
  - (g) -OCF<sub>3</sub>;
  - (h) -CO<sub>2</sub>R<sup>9</sup>,
  - (i) -NR<sup>10</sup>R<sup>11</sup>, and
  - (j) -CONR<sup>10</sup>R<sup>11</sup>;

or a pharmaceutically acceptable salt thereof.

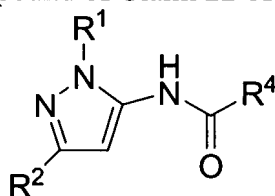
22. (New) The compound of Claim 21 of the formula Ia:



Ia

or a pharmaceutically acceptable salt thereof.

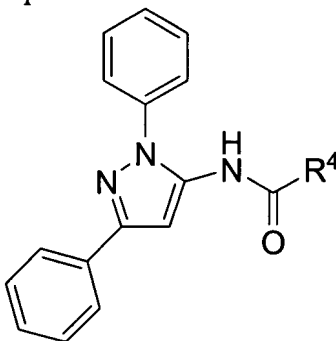
23. (New) The compound of Claim 22 of the formula Ib:



Ib

or a pharmaceutically acceptable salt thereof.

24. (New) The compound of Claim 23 of the formula Ic:



Ic

or a pharmaceutically acceptable salts thereof.

25. (New) The compound of Claim 21 wherein R<sup>1</sup> is phenyl.

26. (New) The compound of Claim 21 wherein R<sup>3</sup> is hydrogen.

27. (New) The compound of Claim 21 wherein  $R^4$  is phenyl, which is unsubstituted or substituted with one or more substituents independently selected from:

- (a)  $-C_{1-6}$ alkyl,
- (b)  $-O-C_{1-6}$ alkyl,
- (c) halo,
- (d) hydroxy,
- (e) trifluoromethyl,
- (f)  $-OCF_3$ ;
- (g)  $-CO_2-C_{1-6}$ alkyl,
- (h)  $-CN$ ,
- (i)  $-NH_2$ ,
- (j)  $-NH-C_{1-6}$ alkyl,
- (k)  $-CONH_2$ , and
- (l)  $-CONH-C_{1-6}$ alkyl.

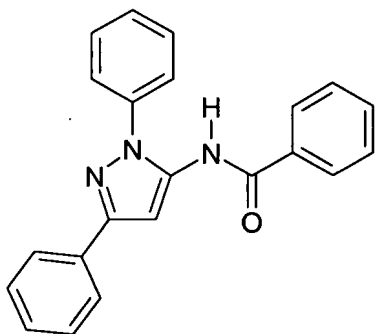
28. (New) The compound of Claim 27 wherein  $R^4$  is phenyl, which is unsubstituted or substituted with halo or  $-CN$ .

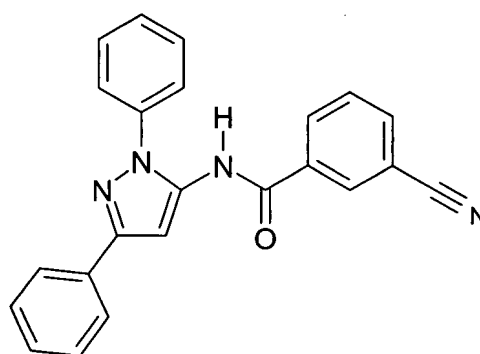
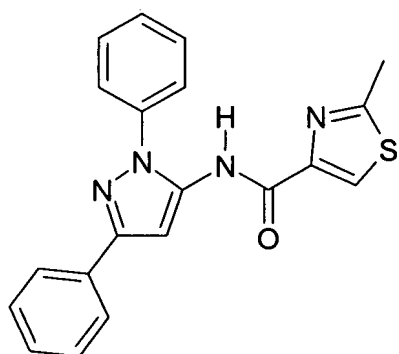
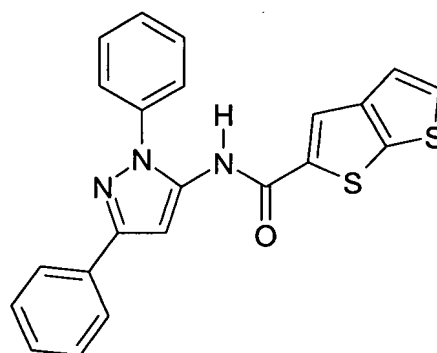
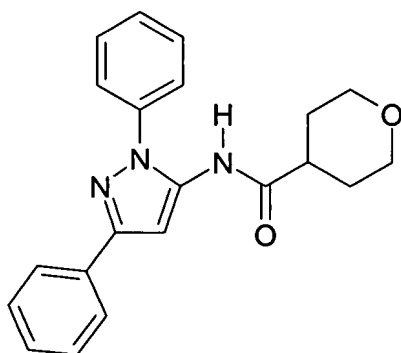
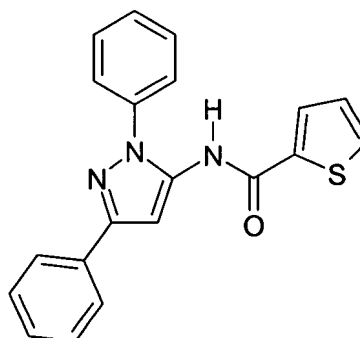
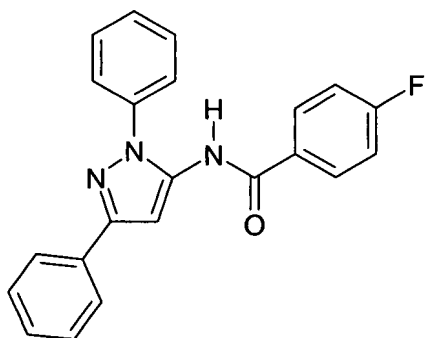
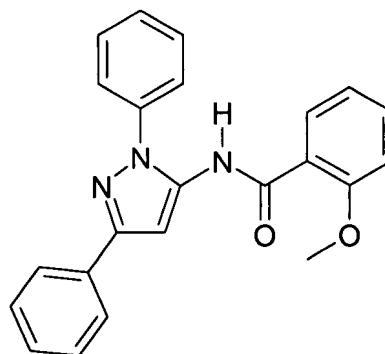
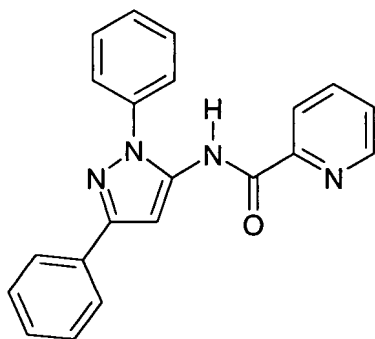
29. (New) The compound of Claim 28 wherein  $R^4$  is phenyl.

30. (New) The compound of Claim 21 wherein  $R^4$  is pyridyl.

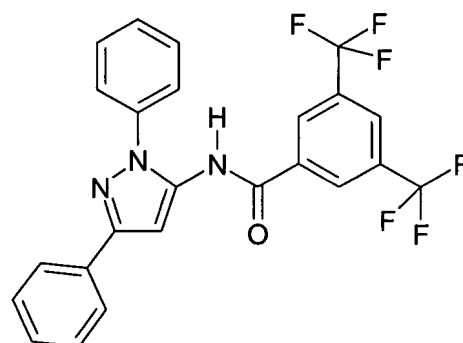
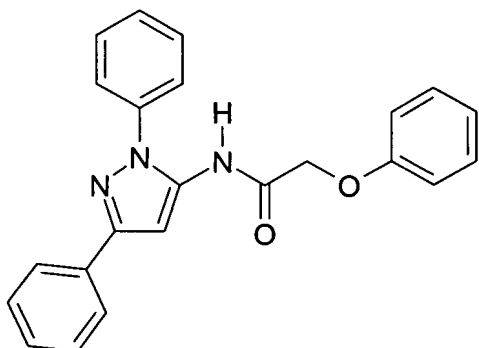
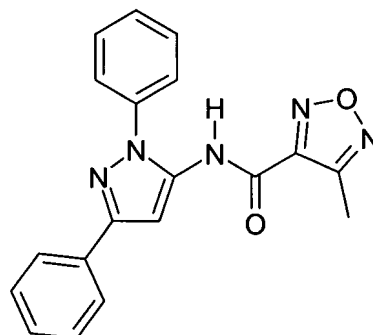
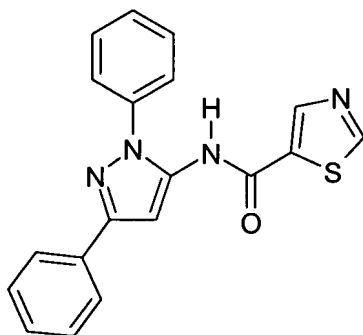
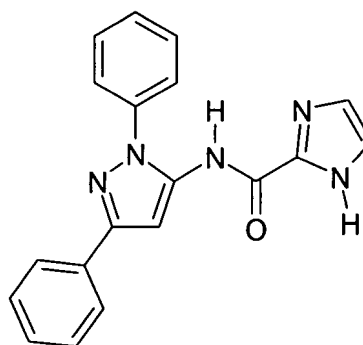
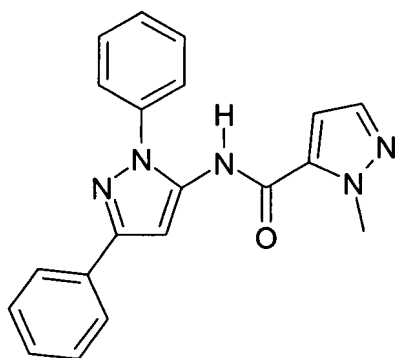
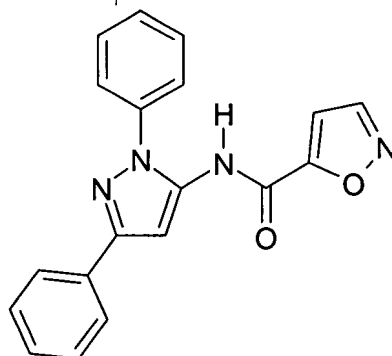
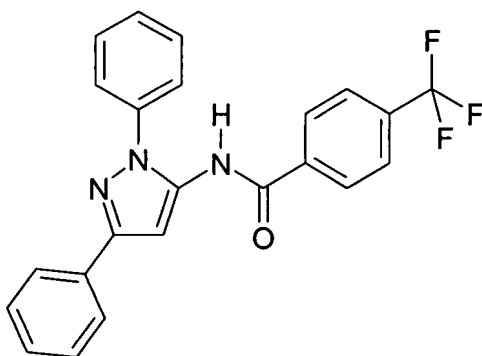
31. (New) The compound of Claim 21 wherein  $R^5$  is hydrogen.

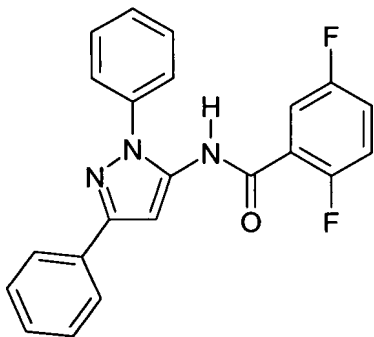
32. (New) A compound which is selected from the group consisting of:











or a pharmaceutically acceptable salt thereof.

33. (New) A pharmaceutical composition which comprises an inert carrier and the compound of Claim 21 or a pharmaceutically acceptable salt thereof.

34. (New) A method for treating schizophrenia in a mammalian patient in need of such which comprises administering to the patient a therapeutically effective amount of the compound of Claim 21 or a pharmaceutically acceptable salt thereof.

35. (New) A method for treating anxiety in a mammalian patient in need of such which comprises administering to the patient a therapeutically effective amount of the compound of Claim 21 or a pharmaceutically acceptable salt thereof.